

ABSTRACT OF THE DISCLOSURE

The present invention relates to tamandarin and didemnin analogs which have a deoxo-proline residue or a dehydro-proline residue in their structure. These analogs are useful as anti-cancer agents and for other purposes. Methods of making
5 these analogs and methods of using them as inhibitors of protein synthesis, cell growth, and tumorigenesis and as enhancers of apoptosis are also provided.

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